

Substitute for form 1449A/PTO

(Use as many sheets as necessary)

Application Number	10/578,105
Filing Date	January 10, 2007
First Named Inventor	Sharma et al.
Art Unit	1654
Examiner Name	Andrew D. Kosar
Attorney Docket Number	1674/2

Sheet	1	of	3	Attorney Docket Number	1674/2
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**INFORMATION DISCLOSURE  
STATEMENT BY APPLICANT**

(Use as many sheets as necessary)

**Complete if Known**

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Sheet 2 of 3

**NON PATENT LITERATURE DOCUMENTS**

Examiner Initials*	Cite No.	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T <sup>2</sup>
	3	Atkinson et al., <u>Peptide Inhibitors of CDK2-cyclin A that Target the Cyclin Recruitment-Site: Structural Variants of the C-Terminal Phe.</u> <u>Bioorganic &amp; Medicinal Chemistry Letters</u> . Vol. 12 pgs.:2501-2505 (2002).	
	4	Boffa et al., <u>Rapamycin Inhibits the Growth and Metastatic Progression of Non-Small Cell Lung Cancer.</u> <u>Clinical Cancer Research</u> . Vol. 10 pgs.:293-300 (2004).	
	5	Brown et al., <u>Rapamycin is active against B-precursor leukemia in vitro and in vivo, an effect that is modulated by IL-7-mediated signaling.</u> <u>PNAS</u> . Vol. 100, No. 25 pgs.:15113-15118 (2003).	
	6	Chen et al., <u>Selective killing of transformed cells by cyclin/cyclin-dependent kinase 2 antagonists.</u> <u>PNAS</u> . Vol. 96 pgs.:4325-4329 (1999).	
	7	Dudkin et al., <u>Biochemical Correlates of mTOR Inhibition by the Rapamycin Ester CCI-779 and Tumor Growth Inhibition.</u> <u>Clinical Cancer Research</u> . Vol. 7 pgs.:1758-1764 (2001).	
	8	Elit, CCI-779 Wyeth. <u>Current Opinion in Investigational Drugs</u> . Vol. 3, No. 8 pgs.:1249-1253 (2002).	
	9	Gali-Muhtasib, H., and Bakkar, Nadine, <u>Modulating Cell Cycle: Current Applications and Prospects for Future Drug Development.</u> <u>Current Cancer Drug Targets</u> . Vol. 2 pgs.:309-336 (2002).	
	10	Huang, S., and Houghton, P.J., <u>Inhibitors of mammalian target of rapamycin as novel antitumor agents: From bench to clinic.</u> <u>Current Opinion in Investigational Drugs</u> . Vol. 3, No. 2 pgs.:295-304 (2002).	
	11	McInnes et al., <u>Peptidomimetic Design of CDK Inhibitors Targeting the Recruitment Site of the Cyclin Subunit.</u> <u>Current Medicinal Chemistry - Anti-Cancer Agents</u> . Vol. 3 pgs.:57-69 (2003).	
	12	Punt et al., <u>Phase I and pharmacokinetic study of CCI-779, a novel cytostatic cell-cycle inhibitor, in combination with 5-fluorouracil and leucovorin in patients with advanced solid tumors.</u> <u>Annals of Oncology</u> . Vol. 14 pgs.:931-937 (2003).	

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	13	Sehgal et al., <i>Rapamycin (AY-22,989), A New Antifungal Antibiotic. II. Fermentation, Isolation and Characterization. The Journal of Antibiotics.</i> Vol. 28, No. 10 pgs.:727-732 (1975).	
	14	Sielecki et al., <i>Cyclin-Dependent Kinase Inhibitors: Useful Targets in Cell Cycle Regulation. Journal of Medicinal Chemistry.</i> Vol. 43, No. 1 pgs.:1-18 (2000).	
	15	Vézina et al., <i>Rapamycin (AY-22,989), A New Antifungal Antibiotic. I. Taxonomy of the Producing Streptomyces and Isolation of the Active Principle. The Journal of Antibiotics.</i> Vol. 28, No. 10 pgs.:721-726 (1975).	
	16	Wiederrecht et al., <i>Mechanism of action of rapamycin: New insights into the regulation of G<sub>1</sub>-phase progression in eukaryotic cells. Progress in Cell Cycle Research.</i> Vol. 1 pgs.:53-71 (1995).	
	17	Zheleva et al., <i>Highly potent p21<sup>WAF1</sup>-derived peptide inhibitors of CDK-mediated pRb phosphorylation: Delineation and structural insight into their interactions with cyclin A. Journal of Peptide Research.</i> Vol. 60 pgs.:257-270 (2002).	

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